

Palynology of the Eocene-Oligocene transition in the marginal zone of the Magura Nappe at Folusz (Western Carpathians, Poland)

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(Manuscript received July 10, 2003; accepted in revised form June 16, 2004)

Abstract: Palynological investigations of the deep-marine Upper Eocene–Lower Oligocene, mainly turbiditic and hemipelagic sediments exposed at Folusz (Siary Zone of Magura Nappe, Polish part of the Western Carpathians), revealed a prevalence of land plant remains in the palynofacies. Dinoflagellate cysts are the most frequent among marine palynomorphs. A Priabonian age is found in the lower part of the Szymbark Shale; a Rupelian age is determined for the investigated part of the Magura Beds. Dinoflagellate cyst distribution shows no major changes, contrary to an outstanding change recorded in the coeval sediments from other parts of the Polish Carpathians. This implies different paleoenvironmental conditions in the northern part of the Magura Basin during the latest Eocene and Early Oligocene. Occurrence of high-latitude dinoflagellate cysts in the lowermost part of the section may be related to a drop of temperature in the surface waters of the sedimentary basin during the Late Eocene. Dinoflagellate cysts, whose motile stages are thought to have inhabited near-shore waters, are frequent in turbidites, whereas the hemipelagic/pelagic sediments usually contain more numerous oceanic specimens. Recycled dinoflagellate cysts occur more frequently in turbidite sediments.

Key words: Carpathians, Eocene–Oligocene transition, Magura Nappe, paleoenvironment, biostratigraphy, flysch deposits, dinocysts.

Introduction

The Eocene–Oligocene transition in the northern part of the Magura Nappe (Siary Zone *sensu* Koszarski et al. 1974) of the Polish Carpathians is represented by a rock succession different from coeval successions of the other parts of the Polish Outer Carpathians. Except for some localities in the eastern sector of the Siary Zone, the succession consists mainly of thick-bedded (0.5 to a few meters) turbidites with variable sandstone to shale ratio, distinguished as the Magura Beds (Szajnocha 1895; Świdziński 1934). In some places, in the marginal part of the eastern sector of the Siary Zone, including the Folusz area (Fig. 1), the Priabonian consists nearly exclusively of fine-grained sediments of turbiditic and hemipelagic origin, distinguished by Kopciowski (1996) as the Szymbark Shale and included in the Zembrzyce Shale by Oszczytko-Clowes (2001). Age-assessments based on foraminifers indicate that the lower boundary of typical Magura Beds (see Książkiewicz 1974) is here located close to the Eocene–Oligocene boundary (Blaicher & Sikora 1963; Sikora 1970; Leszczyński & Malata 2002). At the same time, in the major part of the inner zones of the Magura Nappe, this succession is restricted to the Eocene (Oszczytko-Clowes 2001).

In the other parts of the Polish Outer Carpathians, the Eocene–Oligocene transition is represented by a several meters thick package of *Globigerina*-rich marlstones, distinguished as the Sub-Menilite *Globigerina* Marl (Koszarski & Wieser 1960). The lower part of the overlying unit, known as the Menilite Beds, is dominated by black, organic-rich fine-grained

sediments. Similar sediments are also known from some areas in the southernmost part of the Magura Nappe. The package of marls rich in *Globigerina* is distinguished there as the Lełuchów Marl Member, whereas the unit corresponding to the Menilite Beds is called the Smereczek Shale Member (Birkenmajer & Oszczytko 1989). The so developed Eocene–Oligocene transition displays a major turnover in microfossil assemblages. Rich and diversified Late Eocene foraminifers, calcareous nannoplankton and dinoflagellate cyst assemblages become significantly impoverished in the earliest Oligocene (e.g. Olszewska 1984, 1985; Van Couvering et al. 1981; Gedl 1999). Analysis of organic-walled dinoflagellate cysts (hereafter dinocysts) from these sediments implies that significant paleogeographical and paleoenvironmental changes occurred, including a drop in sea surface temperature and salinity decrease within the photic zone. This paper aims at describing and interpreting the palynological record of the Eocene–Oligocene transition in one section from the northern part of the Magura Nappe, outstanding for its distinctive lithological development. Interpretation of the results is compared with the results of palynological studies of coeval sediments characterized by a markedly different development.

Section location and stratigraphy

A 114 m thick succession representing the Szymbark Shale and the lower part of the Magura Beds, exposed in the bed of Kłopotnica Stream in the village of Folusz was selected for the

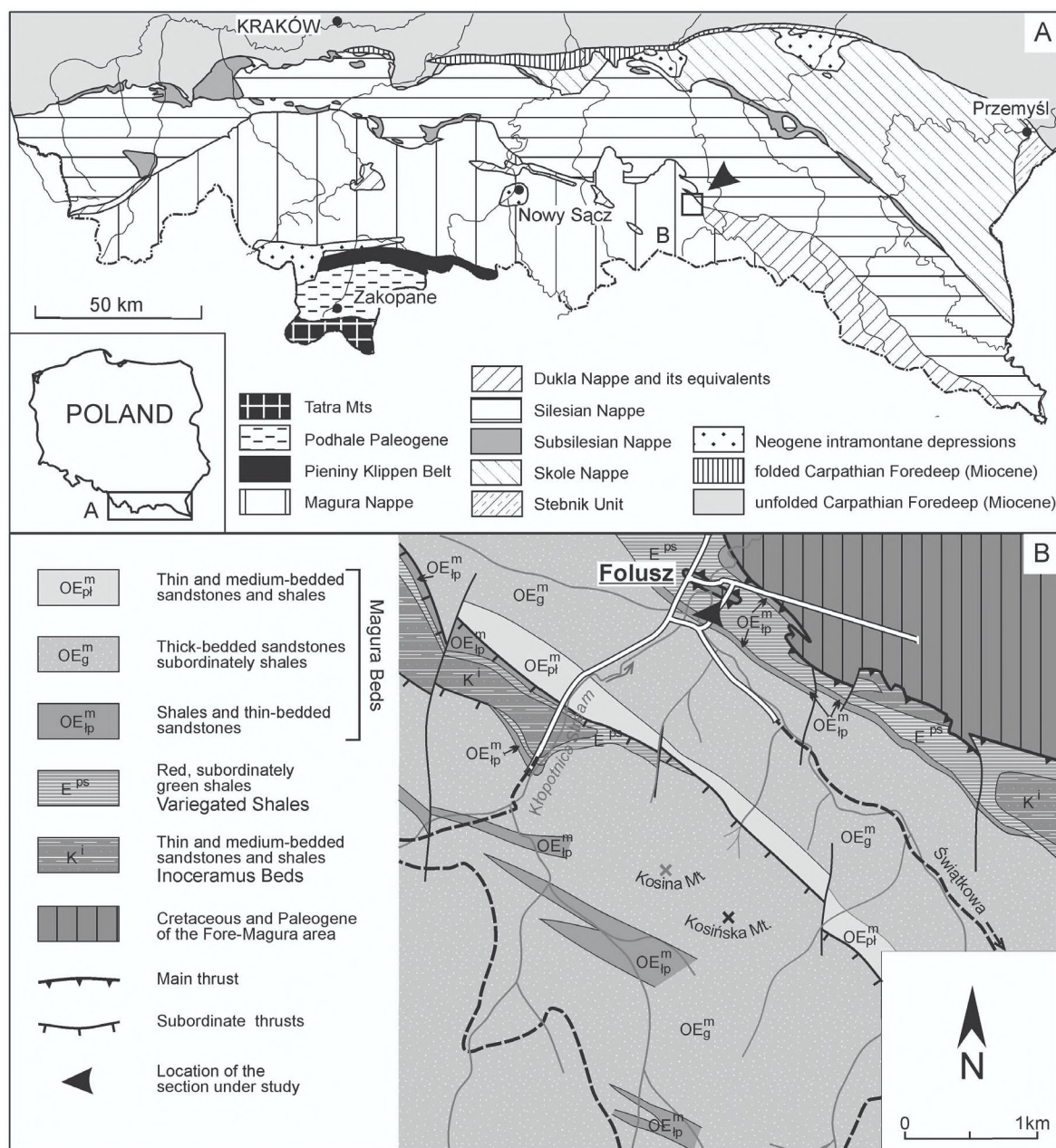


Fig. 1. Location of the studied section (arrowed): **A** — according to tectonic map of the Polish Carpathians after Książkiewicz (1977); **B** — according to geological map after Koszarski & Tokarski (1968). Unit marked OE_{ip}^m in northern outcropping zone includes the Szymbark Shale.

present study (Fig. 1). This section is one of the best-exposed sections of the basal part of the Magura Beds and their immediate substrate in the frontal part of the Magura Nappe (Fig. 1A). This section is also best recognized in the sense of litho- and biostratigraphy (see Sikora 1970; Oszczytko-Clowes 2001; Leszczyński & Malata 2002). Moreover, tuffite recorded here was dated radiometrically (Van Couvering et al. 1981).

The Folusz section is located in the central part of the Polish Carpathians, at the northern foots of the Magura Wątkowska Range in the Beskid Niski Mountains. The section starts at the upper boundary of the red shale succession (Łabowa Shale Formation *sensu* Oszczytko 1991) and extends up the stream, up to the bridge of a local road toward the village of

Świątkowa (Fig. 1B). The lower part of the section, approximately 36 meters thick, called by Kopciowski (1996) the Szymbark Shale, consists basically of alternating thin layers of light green to dark green, occasionally black and dark brown muddy to clayey, calcareous to non-calcareous shales (Fig. 2). The proportion of calcareous and dark coloured shales increases up section. Moreover, several thin siltstone and fine-grained sandstone beds, and one 4 cm thick layer of bentonitic tuffite occur in this unit. The position of the tuffite in the section (Fig. 2), together with its thickness and colour, suggest that this is the tuffite mentioned earlier by Sikora (1970). Zircons from this layer were dated by fission-track analysis at 32.8 ± 1.3 Ma (Van Couvering et al. 1981), i.e. close to the

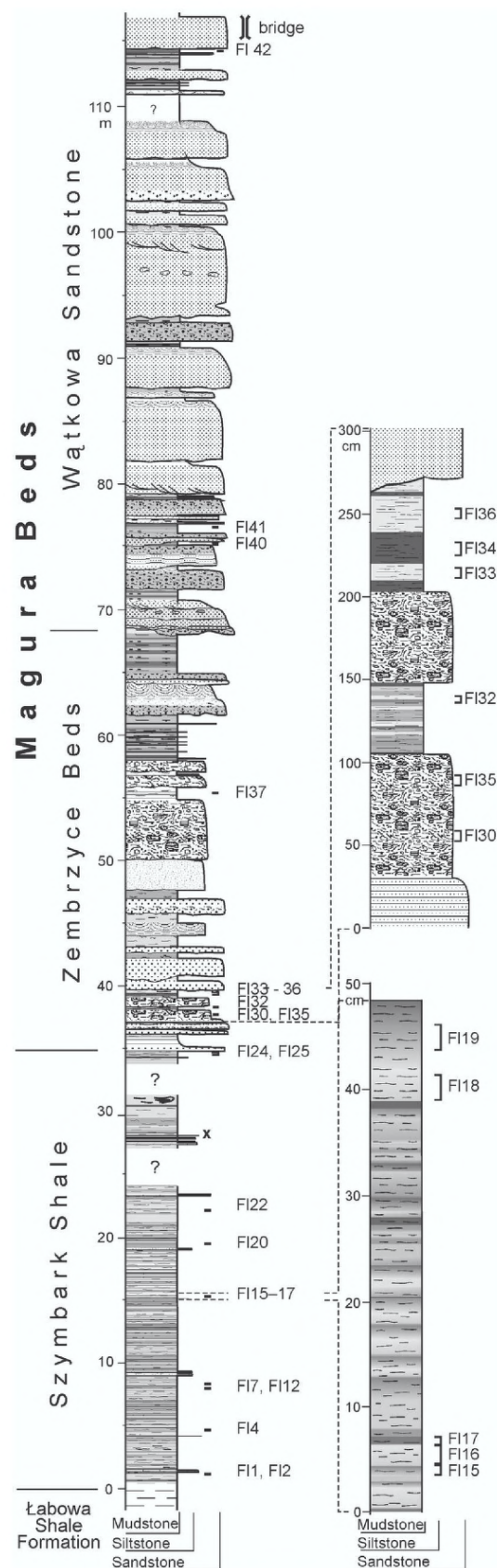


Fig. 2. Vertical lithofacies logs and lithostratigraphy of the studied section at Folusz, and sample locations (cf. Leszczyński & Malata 2002).

Eocene-Oligocene boundary (33.7 Ma according to Berggren et al. 1995). In the upper part of this unit, Blaicher & Sikora (1963) found a foraminiferal assemblage corresponding to that recorded in the Sub-Menilite Globigerina Marl (upper part of Upper Eocene) of the outer flysch nappes. These authors called this interval "Marls with Globigerina" (cf. Sikora 1970). The calcareous nannoplankton Zone NP19-20 (Martini 1971) was recognized in the lower part of the Szybark Shale, and Zone NP21 in its upper part, ca. 11 m above the base of the unit in Oszczytko-Clowes (2001), or 17.0 m in Leszczyński & Malata (2002).

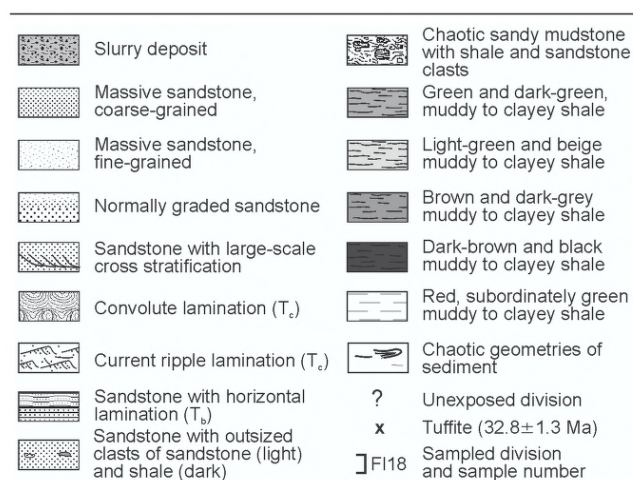
The overlying 33 m thick succession is built of thin to thick-bedded fine to very coarse-grained sandstones, chaotic sandy-mudstones and thick to thin layers of calcareous to non-calcareous mudstones (Fig. 2). This unit is called here the Zembrzyce Beds, and considered to be the basal part of the Magura Beds. Calcareous nannoplankton investigations indicate that the entire unit is enclosed within Zone NP21.

Very thick-bedded coarse- to fine-grained sandstones and granule-conglomerates dominate in the remaining part of the section (Fig. 2). This part of the section is considered as the Wątkowa Sandstone *sensu* Koszarski (1976). It represents the higher part of the Magura Beds. The base of calcareous nannoplankton Zone NP22 was identified in the top part of the section (Oszczytko-Clowes 2001).

Materials

Twenty-one samples from surface outcrops in the Kłopotnica Stream were investigated in this study. These samples were selected from those taken earlier for other sedimentological and stratigraphical investigations (Leszczyński & Malata 2002).

Eleven samples (FI1-FI25) represent the Szybark Shale, seven samples (FI30-FI37) were taken from the Zembrzyce Beds, and three (FI40-FI42) from the Wątkowa Sandstone (Fig. 2). All samples except for the FI30 and FI35 were taken from muddy to clayey shales. Their location in the sequence as well as foraminiferal content and composition suggest



Legend to the Fig. 2.

Inferred age (this study)	Late Eocene	Late Eocene- Early Oligocene		Early Oligocene																		
Lithostratigraphy	Szymbark Shale												Magura Beds									
													Zembrzyce Beds							Wątkowa Sandstone		
Taxon \ Sample	F1	F2	F4	F7	F12	F15	F16	F17	F20	F22	F24	F25	F30	F35	F32	F33	F34	F36	F37	F40	F41	F42
<i>Spiniferites ramosus</i>																						
<i>Phthanoperidinium comatum</i>																						
<i>Rhombodinium perforatum</i>																						
<i>Homotryblum tenuispinosum</i>																						
<i>Lentinia serrata</i>																						
<i>Areosphaeridium michoudii</i>																						
<i>Areosphaeridium diktyoplokum</i>																						
<i>Selenopemphix nephroides</i>																						
<i>Deflandrea</i> spp.																						
<i>Selenopemphix coronata</i>																						
<i>Cerebrocysta bartonensis</i>																						
<i>Corrudinium incompositum</i>																						
<i>Operculodinium centrocarpum</i>																						
<i>Pentadinium laticinctum</i>																						
<i>Impagidinium disperitum</i>																						
<i>Impagidinium aculeatum</i>																						
<i>Dracodinium laszczyński</i>																						
<i>Impagidinium</i> sp. A																						
<i>Impagidinium</i> sp. B																						
<i>Gelatia inflata</i>																						
<i>Polysphaeridium subtile</i>																						
<i>Pentadinium laticinctum granulosum</i>																						
<i>Spiniferites pseudofurcatus</i>																						
<i>Rhombodinium freienwaldensis</i>																						
<i>Impagidinium velorum</i>																						
<i>Cordosphaeridium funiculatum</i>																						
<i>Phelodinium</i> sp.																						
<i>Lejeunecysta tenella</i>																						
<i>Spiniferites membranaceus</i>																						
<i>Operculodinium microtriainum</i>																						
<i>Lingulodinium machaerophorum</i>																						
<i>Membranophoridium aspinatum</i>																						
<i>Impagidinium brevisculatum</i>																						
<i>Homotryblum vallum</i>																						
<i>Trigonopyxidia ginella</i>																						
<i>Leptodinium membranigerum</i>																						
<i>Tectatodinium</i> aff. <i>pellitum</i>																						
<i>Homotryblum aculeatum</i>																						
<i>Distatodinium ellipticum</i>																						
<i>Nannoceratopsis gracilis</i>																						
<i>Nematosphaeropsis labyrinthus</i>																						
<i>Selenopemphix</i> aff. <i>selenoides</i>																						
<i>Impagidinium pallidum</i>																						
<i>Melittasphaeridium pseudorecurvatum</i>																						
<i>Hystrichokolpoma rigaudiae</i>																						
<i>Palaeocystodinium golzowense</i>																						
<i>Homotryblum plectilum</i>																						
<i>Lejeunecysta</i> sp.																						
<i>Achomosphaera ramulifera</i>																						
<i>Systematophora placacantha</i>																						
<i>Tityrosphaeridium cantharellus</i>																						
<i>Areoligera undulata</i>																						
<i>Enneadocysta pectiniformis</i>																						
<i>Charlesdowniea coleothrypta</i>																						
<i>Enneadocysta</i> aff. <i>pectiniformis</i>																						
<i>Glaphyrocysta semitecta</i>																						
<i>Lejeunecysta fallax</i>																						

Fig. 3. Range chart of dinocysts from Folusz section. Line widths reflect number of specimens counted. Continued on the page 159.

hemipelagic and/or turbiditic origin (Table 1; Leszczyński & Malata 2002). The samples F130 and F135 were taken from a chaotic sandy-mudstone breccia. This is a sediment consisting of a sandy calcareous mudstone (marlstone) matrix with quartz grains as much as 7 mm in cross-section and chaotically distributed dark green, brownish and grey, sharp-edged mudstone chips and larger clasts. These features indicate deposition by debris flow mechanisms.

Methods

The samples were processed following standard palynological procedure, including 38% hydrochloric acid (HCl) treatment, 40% hydrofluoric acid (HF) treatment, heavy liquid ($\text{ZnCl}_2 + \text{HCl}$; density 2.0 g/cm^3) separation, ultrasound for 10–15 sec., and sieving at $15 \mu\text{m}$ on a nylon mesh. The samples were processed in the Micropaleontological Laboratory of the Institute of Geological Sciences, Polish Academy of Sciences, Kraków. The quantity of rock processed varied from 20 to 30 g. Two microscope slides were made from each sample using glycerine jelly as a mounting medium. The rock samples, palynological residues and slides are stored in the collection of the Institute of Geological Sciences, Polish Academy of Sciences, Kraków.

All dinocysts were counted from one of two slides. The second slide was scanned for additional dinocyst taxa. Slides were examined under the transmitted light microscope Carl Zeiss Axiolab. Microphotographs were taken with the Sony DSC-S75 camera and Zeiss Plan-Neofluar $100 \times/1.30$ Oil Pol objective.

Results

All samples contain organic particles of a diameter greater than $15 \mu\text{m}$. The majority of these particles represent land plant remains. They include cuticle with variously preserved tissue structures, dark brown equidimensional particles with slightly transparent edges and opaque, small-sized, woody particles. Sporomorphs (mostly the bisaccate pollen grains) occur subordinately. Amorphous organic matter occurs in trace amounts only.

Marine palynomorphs, rarely exceeding 1–2 % of the palynofacies, are represented primarily by dinocysts (Figs. 3, 4). Acritarchs and foraminiferal linings occur occasionally. The highest amount of dinocysts, reaching 8 % of the palynofacies, is recorded in sam-

Inferred age (this study)	Late Eocene	Late Eocene- Early Oligocene	Early Oligocene																			
Lithostratigraphy	Szymbark Shale										Magura Beds											
											Zembrzyce Beds								Wątkowa Sandstone			
Taxon	Sample	F11	F12	F14	F17	F112	F115	F116	F117	F120	F122	F124	F125	F130	F132	F133	F134	F136	F137	F140	F141	F142
<i>Melitasphaeridium asterium</i>																						
<i>Glaphyrocysta</i> sp. A																						
<i>Reticulatosphaera actinocoronata</i>																						
<i>Operculodinium</i> aff. <i>centrocarpum</i>																						
<i>Ctenidodinium omatum</i> /combazii																						
<i>Adnatosphaeridium vittatum</i>																						
<i>Wetzeliiella symmetrica</i>																						
<i>Rottnestia borussica</i>																						
<i>Cribroperidinium tenuitabulatum</i>																						
<i>Heteraulacacysta porosa</i>																						
<i>Amphorosphaeridium?</i> <i>multispinosum</i>																						
<i>Dinopterygium</i> sp. A																						
<i>Selenopemphix armata</i>																						
<i>Fibrocysta</i> aff. <i>vectensis</i>																						
<i>Lejeunecysta hyalina</i>																						
<i>Enneadocysta arcuata</i>																						
<i>Surculosphaeridium?</i> <i>longifurcatum</i>																						
<i>Hystriocholopoma cinctum</i>																						
<i>Fibrocysta vectensis</i>																						
<i>Charlesdownia</i> sp.																						
<i>Operculodinium deconinckii</i>																						
<i>Impagidinium gibrense</i>																						
<i>Diphyes colligerum</i>																						
<i>Rottnestia</i> aff. <i>borussica</i>																						
<i>Chatangiella ditissima</i>																						
<i>Impagidinium</i> sp. C																						
<i>Lingulodinium pycnospinosum</i>																						
<i>Thalassiphora</i> sp. A																						
<i>Areoligera</i> sp.																						
<i>Diphyes pseudoficusoides</i>																						
<i>Wilsonidinium intermedium</i>																						
<i>Wetzeliiella gochti</i>																						
<i>Samlandia chlamydochora</i>																						
<i>Pyxidinospis</i> sp. A																						
<i>Cyclonephelium compactum</i>																						
<i>Operculodinium?</i> <i>hirsutum</i>																						
<i>Corrudinium?</i> sp. A																						
<i>Pseudoceratium pelliferum</i>																						
<i>Pyxidinospis?</i> sp. B																						
<i>Cordosphaeridium inodes</i>																						
<i>Areoligera semicirculata</i>																						
<i>Homotryblium?</i> sp. A																						
<i>Paucisphaeridium inversibuccinum</i>																						
<i>Pyxidinospis</i> sp. C																						
<i>Eatonocysta ursulae</i>																						
<i>Polysphaeridium zoharyi</i>																						
<i>Glaphyrocysta intricata</i>																						
<i>Heteraulacacysta</i> sp. A																						
<i>Pyxidinospis psilata</i>																						
<i>Adnatosphaeridium multispinosum</i>																						
<i>Phthanopendinium</i> aff. <i>amiculum</i>																						
<i>Cordosphaeridium minimum</i>																						
<i>Achomosphaera alcornu</i>																						
<i>Selenopemphix</i> sp. A																						
<i>Dapsilidinium multispinosum</i>																						
<i>Pyxidinospis</i> sp. D																						
<i>Phthanopendinium amiculum</i>																						
<i>Pyxidinospis</i> sp. E																						
<i>Rhombodinium draco</i>																						
<i>Hystriochosphaeridium</i> sp.																						
<i>Glaphyrocysta</i> sp. B																						
<i>Lejeunecysta lata</i>																						
<i>Rhombodinium</i> sp. A																						
<i>Florentinia</i> sp.																						
<i>Impagidinium</i> sp. D																						
<i>Thalassiphora</i> aff. <i>pelagica</i>																						
<i>Glaphyrocysta</i> sp.																						
<i>Pentadinium?</i> <i>circumsutum</i>																						
<i>Dinopterygium</i> sp. B																						
<i>Adnatosphaeridium</i> sp. A																						
<i>Nannoceratopsis dictyambonis</i>																						
<i>Achomosphaera sagena</i>																						
<i>Turbiosphaera symmetrica</i>																						
<i>Apectodinium quinquelatum</i>																						
<i>Enneadocysta multicornuta</i>																						
<i>Dapsilidinium pastielsii</i>																						
<i>Diphyes</i> aff. <i>colligerum</i>																						

ples F14 and F17, representing the Szymbark Shale. Samples F134 and F141 (the Zembrzyce Beds and the Wątkowa Sandstone respectively) do not contain dinocysts or contain only a few indeterminate specimens. Acritarchs occur mainly in the lower part of the Szymbark Shale (samples F11–F17), being the most frequent in sample F14. They represent morphotypes of *Veryhachium* affinity. Microforaminifers (mainly trochospiral and planispiral morphotypes) are very rare within the whole section except for the sample F14 where they are relatively frequent.

The dinocyst distribution shows distinct relation to the sediment origin (Fig. 5). Samples taken from turbidites and mud-flow deposits (Table 1) are either devoid of oceanic dinocysts (e.g. *Impagidinium* spp.; see e.g. Dale 1996), or they occur there as single specimens only. Specimens of *Impagidinium* (Fig. 4.3,6,8,13) are much more frequent in samples representing hemipelagites (Table 1). In these samples, *Impagidinium* constitutes up to 30 % of all dinocysts (sample F122). The samples taken from turbidites, in turn, contain high numbers of *Homotryblium* spp., a genus related by several authors to inshore, lagoonal settings (e.g. Brinkhuis 1994).

Dinocyst stratigraphy of the Folusz section

The occurrence of calcareous foraminifers and nannoplankton in the studied succession mainly in resedimented deposits (see Leszczyński & Malata 2002) hampers a precise interpretation of stratigraphy based on these fossils. Recycling also concerns dinocysts as has been shown in these investigations. However, dinocysts also occur in background sediments (hemipelagites) where they appear to be syndimentary. Re-working makes particularly tentative the identification of biozones or events defined according to the highest occurrences of fossils, for example the NP21 and the NP22 Calcareous Nannoplankton Zones of Martini (1971).

Location of the Eocene-Oligocene boundary in the investigated section has not been precisely established during this study. The Early Oligocene has been definitely determined for the section division located above sample F124 (the topmost part of the Szymbark Shale) where *Areoligera? semicirculata* (Fig. 4.2) has its lowest occurrence (Fig. 6). This species is known to have its first appearance in the earliest Oligocene (e.g. Morgenroth 1966; Brinkhuis 1994), Rupelian (Stover & Hardenbol 1993; Stover et al. 1996) or Chattian (Powell 1992). This agrees with the age 32.8 ± 1.3 Ma, obtained by Van Couvering et al. (1981) from radiometric dating of the tuffite found approximately 7 m below the sample F124. The age of the Magura Beds, overlying the Szym-

Table 1: Selected sedimentological and geochemical features of the sampled sediments.

Sample No.	Sediment colour	Reaction with HCl	Bed thickness [cm]	CaCO ₃ [%]	TOC [%]	Sand fraction			Sediment origin
						Detritus %	Foraminifera number		
							Agglut.	Calc.	
F11	dark brown	+	3.0	nd	nd	nd	nd	nd	Hemipelagite?
F12	green	—	8.0	nd	nd	nd	nd	nd	Hemipelagite?
F14	brown-grey	+	5.0	17.70	0.46	0.175	572	3256	Turbidite
F17	brown	+	16.0	21.98	0.50	s	3	43	Hemipelagite?
F112	green	—	15.0	0.42	0.18	s	678	1	Hemipelagite
F115	green	—	1.0	nd	nd	s	3208	128	Hemipelagite?
F116	beige	+	2.0	10.25	0.30	s	504	48	Hemipelagite?
F117	dark brown	—	0.7	1.17	1.74	tr	200	18	Hemipelagite?
F120	brown	+	27.0	18.67	0.55	—	1	2	Turbidite
F122	beige and green	+, —	30.0	6.17	0.28	tr	1132	16	Hemipelagite?
F124	grey-green	+	>20	nd	nd	~5.0	44	1190	Turbidite/Hemipelagite
F125	dark brown	—	7.0	0.33	3.20	9.5	64	32	Turbidite
F130	green/dark grey	+	75.0	nd	nd	~12.0	400	20868	Mud-flow deposit
F132	green/dark grey	—, (+)	40.0	nd	nd		68	56	Turbidite/Hemipelagite
F133	grey	—	10.0	nd	nd	tr	22	—	Hemipelagite?
F134	dark brown	—	19.0	nd	nd	tr	—	—	Turbidite
F135	grey/green	+	75.0	nd	nd	~15.0	nd	nd	Mud-flow deposit
F136	grey-green	—, (+)	22.0	nd	nd	tr	8	32	Turbidite/Hemipelagite
F137	grey	+	10.0	nd	nd	tr	22	8	Turbidite
F140	grey-green	+	7.0	nd	nd	nd	nd	nd	Turbidite?
F141	black	—	7.0	nd	nd	nd	nd	nd	Hemipelagite?
F142	grey	+	6.0	nd	nd	nd	nd	nd	Turbidite?

Number of foraminiferal specimens calculated for 200 g of dry rock. See Fig. 2 for sample location. Explanation of symbols: + — effervescent reaction with diluted HCl; (+) — weak effervescence; — — lack of effervescence; tr — <0.1 %; s — several grains; nd — not determined.

bark Shale, can be estimated on the basis of superposition only. No typical Oligocene dinocysts like *Wetzeliiella gochtii*, *Areoligera? semicirculata* or *Chiropteridium lobospinosum* have been found here. The lack of the latter mentioned species may suggest an age not younger than NP23. Powell (1992) and Wilpschaar et al. (1996) recorded the lowest occurrence of *Chiropteridium lobospinosum* within this zone. This species was found in the Polish Carpathians in the Lower Oligocene of the Podhale Flysch (Gedl 2000) and in the Oligocene Krosno Beds, whereas it was not found so far in the lowermost Oligocene in the Menilite Beds (P. Gedl unpubl.).

Other dinocyst events, which may point at the Oligocene age, are rather disputable. One specimen of *Wetzeliiella gochtii* (Fig. 4.20) has been found in sample F120 (Fig. 6). The lowest occurrence of this species is often accepted as an indicator of the Lower Oligocene (e.g. Costa & Downie 1976; Gruas-Cavagnetto & Barbin 1988), although some authors claimed that this species appeared in the Late Eocene (Châteauneuf 1980; El-Beialy 1988). In the Polish Carpathians, specimens of *Wetzeliiella gochtii* have been found in higher part of the Sub-Menilite Globigerina Marl at Krosno and Znamirów (Bujak in Van Couvering 1981) or its equivalent at Leluchów (Gedl 2004). In the latter case, the lowest occurrence of *Wetzeliiella gochtii* comes from a sample where *Areosphaeridium diktyoplokum* has its highest occurrence.

The lowest occurrence of *Reticulatosphaera actinocoronata* (Fig. 4.15), a species believed to have appeared for the first time in the Oligocene in the middle latitudes (e.g. Manum et al. 1989), was recorded in the Priabonian part of the Szymbark Shale (sample F14). On the other hand, Brinkhuis (1994) demonstrated the first appearance of this species in central and northern Italy in the Late Eocene (basal part of the NP21

Zone). An even younger first appearance, early Late Eocene, was recorded by Coccioni et al. (2000) in central Italy. This event was correlated with subChron16n and calcareous nanoplankton Zone NP19/20. The lowest occurrence of *Reticulatosphaera actinocoronata* in the Folusz section can also be correlated with the Zone NP19/20 since this species occurs together with *Areosphaeridium michoudii*.

The age of the lower part of the Szymbark Shale (below the sample F17) is Late Eocene (Priabonian). This is shown by the co-occurrence of dinocyst species *Areosphaeridium diktyoplokum* (Fig. 4.11), *Areosphaeridium michoudii* (Fig. 4.10) and *Rhombodinium perforatum* (Fig. 4.24) in this interval (Fig. 6). The last appearance of *Areosphaeridium diktyoplokum*, which has the highest consistent occurrence in sample F17 (Fig. 6), was commonly believed to have taken place at the Eocene-Oligocene boundary (e.g. Biffi & Manum 1988; Stover et al. 1996), although some authors also found it in the Lower Oligocene (e.g. Maier 1959; Benedek 1986). Brinkhuis (1994) demonstrated the Early Oligocene last appearance of this species in Italy (for discussion see also Berggren et al. 1995 and Brinkhuis & Visscher 1995). *Areosphaeridium michoudii* has its highest known occurrence at the top of the NP18 Zone in the North Sea (Bujak & Mudge 1994). However, distribution of this species in the flysch Carpathians suggests its highest occurrence within the NP19-20 Zone (lower part of the Sub-Menilite Globigerina Marl; P. Gedl unpubl.). The highest consistent occurrence of *Areosphaeridium michoudii* in the studied section is recorded in sample F112, which is located higher than the highest consistent occurrence of *Areosphaeridium diktyoplokum* (sample F17). A Priabonian age of the lower part of the Szymbark Shale in the Folusz section is also indicated by the presence of *Rhombodinium perforatum*. This species

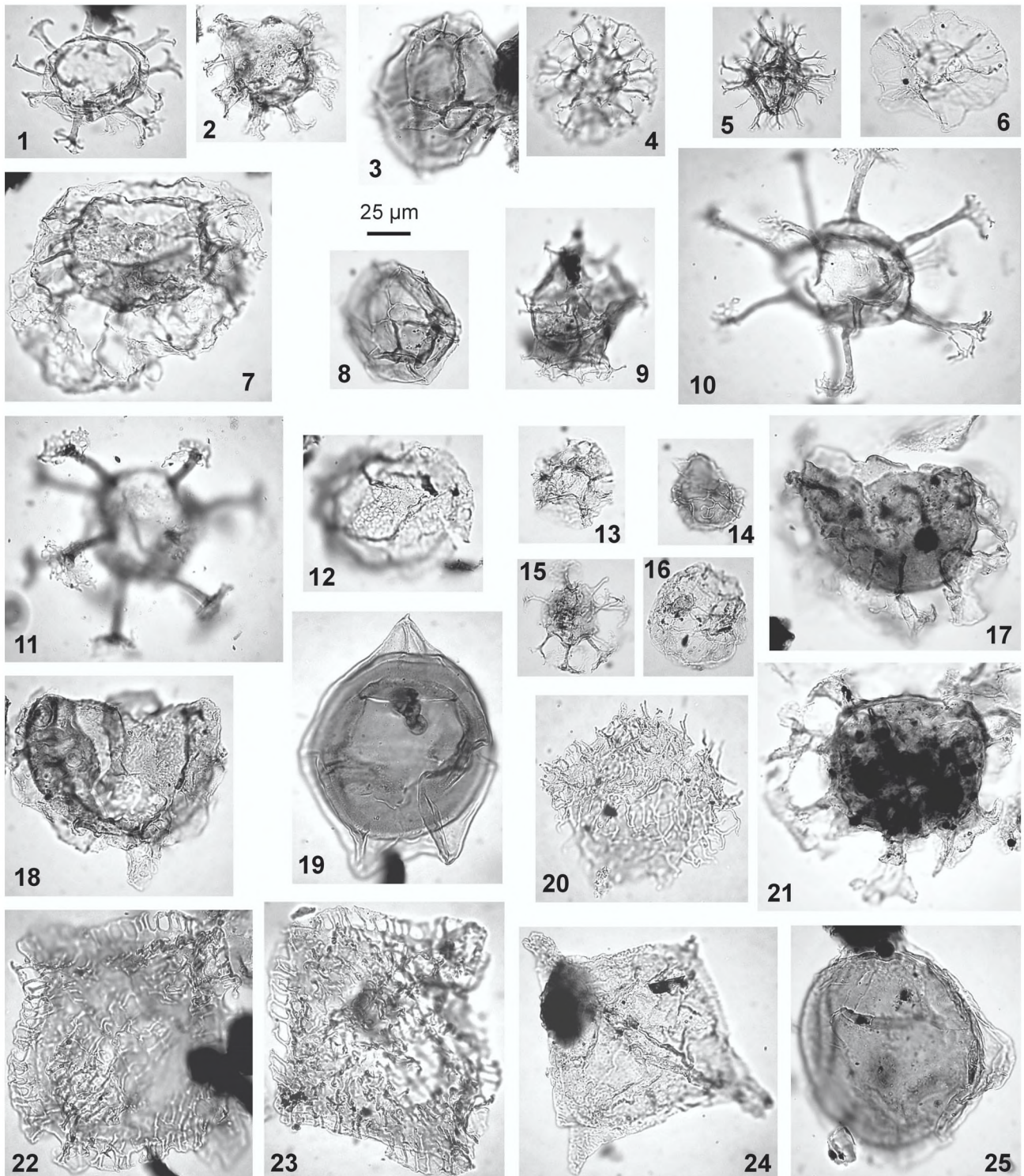


Fig. 4. Dinocysts from the studied section. Scale bar (25 µm) refers to all photomicrographs. Slide code and England Finder references are given. 1 — *Homotryblium plectilum*, Fl35a[H34.1]; 2 — *Areoligera? semicirculata*, Fl33b[E35.2]; 3 — *Impagidinium brevisulcatum*, Fl7[G47.4]; 4 — *Nematosphaeropsis labyrinthus*, Fl2b[J36.2-4]; 5 — *Spiniferites ramosus*, Fl33b[W46.2]; 6 — *Impagidinium velorum*, Fl2a[Q32.2]; 7 — *Glaphyrocysta semitecta*, Fl4a[U36.2]; 8 — *Impagidinium dispertitum*, Fl12a[R48]; 9 — *Rottnestia borussica*, Fl4a[T32.2]; 10 — *Areosphaeridium michoudii*, Fl2a[S45]; 11 — *Areosphaeridium diktyoplokum*, Fl2b[Q40.4]; 12 — *Heterelaucacysta porosa*, Fl4a[V43.1-3]; 13 — *Impagidinium pallidum*, Fl4a[X39.1]; 14 — *Corrudinium incompositum*, Fl12a[R39]; 15 — *Reticulosphaera actinocoronata*, Fl12a[D47.2-4]; 16 — *Gelatia inflata*, Fl1[H37.2]; 17 — *Areoligera undulata*, Fl4a[N30.2]; 18 — *Membranophoridium aspinatum*, Fl4b[W31.1-3]; 19 — *Deflandrea* sp., Fl15b[F36]; 20 — *Wetzeliella gochtii*, Fl20b[C42]; 21 — *Areoligera sentosa*, Fl4b[X47.4]; 22 — *Charlesdowniea clathrata*, Fl4a[T34.1-2]; 23 — *Charlesdowniea coleothrypta*, Fl4b[H46.3]; 24 — *Rhombodinium perforatum*, Fl2b[W35.2-4]; 25 — *Dracodinium laszczynskii*, Fl1[G44.1-3].

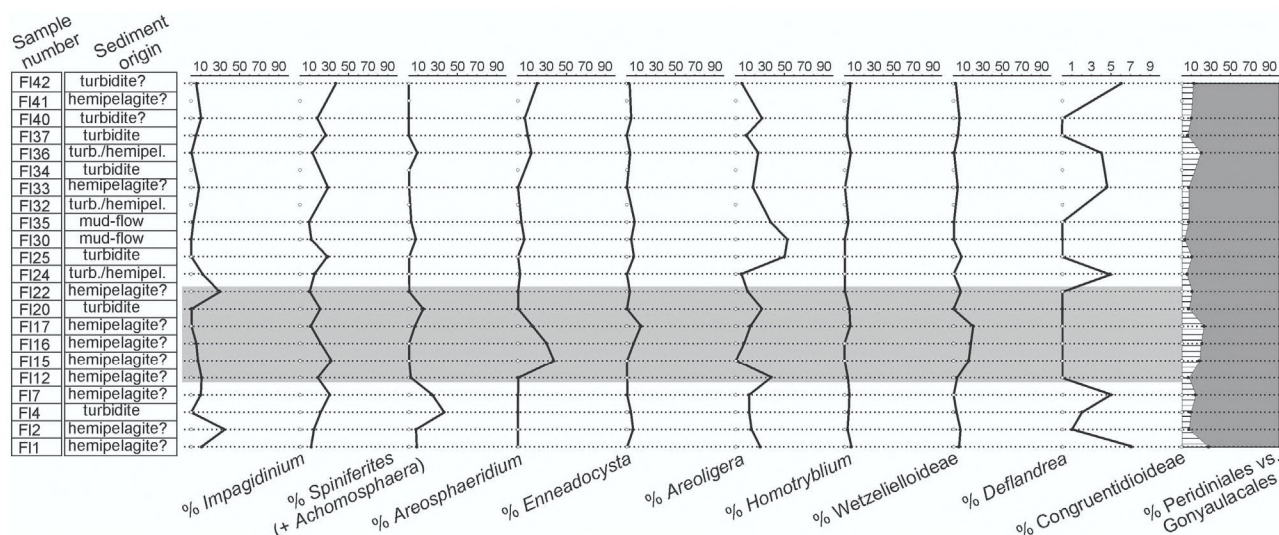


Fig. 5. Frequency of selected dinocysts in relation to sediment origin. Shaded area shows undivided Upper Eocene-Lower Oligocene interval based on dinocysts.

has the lowest occurrence at the base of the NP18 Zone, and ranges to the end of the Eocene (Powell 1992). The age of the interval between sample F112 and F124 has not been precisely documented. It has been assigned as undivided latest Eocene-earliest Oligocene (Fig. 6).

Interpretation of age based on dinocysts agrees in general with that based on foraminifers (Malata *in* Leszczyński &

Malata 2002) and calcareous nannoplankton (Oszczypko-Clowes 2001; Oszczypko-Clowes *in* Leszczyński & Malata 2002) for the lowermost part of the section (Fig. 7). All methods indicate a Late Eocene age of the basal part of the Szymbark Shale.

Significant differences appear in the biostratigraphy of the higher part of the section. Dinocysts indicate a Late Eocene

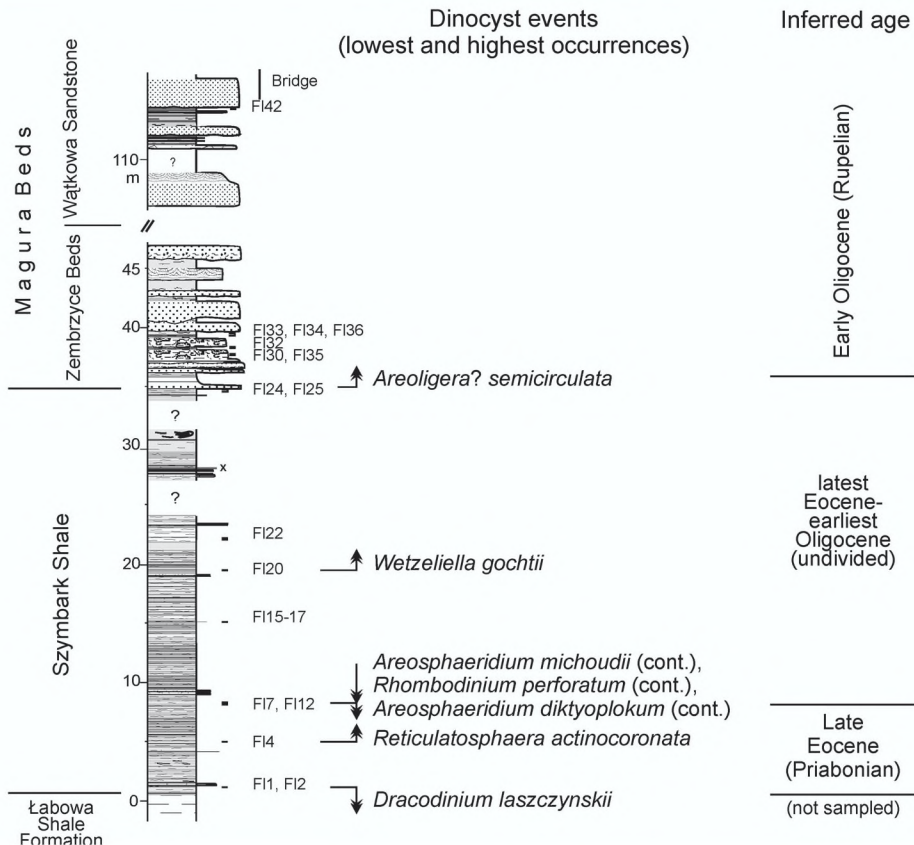


Fig. 6. Lowest and highest occurrences of selected dinocyst species in studied section and age interpretation.

age of the lower part of the Szymbark Shale up to the sample FI7 (basal 8 meters of the section). Foraminifers interpreted by Malata (*in* Leszczyński & Malata 2002) indicate a Late Eocene age up to the sample FI22. The assemblage from samples FI18–FI22 indicates an age close to the Eocene-Oligocene boundary. This means that the lowest occurrence of *Wetzelietta gochtii*, found in sample FI20, is in the uppermost Eocene. Moreover, this implies that *Wetzelietta gochtii* in the Carpathian Flysch Basin appeared for the first time in the latest Eocene. Van Mourik & Brinkhuis (2000) noted a Late Eocene first appearance of this species from a low-latitude setting in the Atlantic Ocean. These records suggest diachronous first appearance of *Wetzelietta gochtii*, starting from low latitudes.

Calcareous nannoplankton, in turn, was interpreted by Oszcypko-Clowes (2001) to indicate the middle Priabonian, corresponding to the Zone NP19–20, for the ca. 11 m thick, basal part of the Szymbark Shale (i.e. approximately up to the position of our sample FI12). Upper Priabonian to lower Rupelian, corresponding to the Zone NP21, was suggested there for the overlying part of the section, which according to the lithostratigraphy used in this paper, embraces the higher part of the Szymbark Shale and majority of the Zembrzyce Beds. Interpretations of calcareous nannoplankton by Oszcypko-Clowes (*in* Leszczyński & Malata 2002) suggest a middle Priabonian age for the Szymbark Shale up to the sample FI18, found at least up to ca. 17 m above the lower boundary of the unit.

Dinocysts indicate the Lower Oligocene (Rupelian) in the section above sample FI24. The age based on foraminifers from sample FI24 was interpreted as undivided Priabonian-Rupelian. The lowest occurrence of Early Oligocene foraminifers was identified first in the sample FI30 (mud-flow deposits containing a rich dinocyst assemblage interpreted as recycled Eocene). This sample is located in the lower part of the Zembrzyce Beds. A higher part of the lower Rupelian, corresponding to the Zone NP22, was identified with calcareous nannoplankton in the topmost part of the Zembrzyce Beds and in the remaining part of the section (Oszczypko-Clowes 2001; note: the position of the NP21 has been erroneously indicated there in fig. 44). Noteworthy, the chronostratigraphy of the succession in question as interpreted by Oszcypko-Clowes (2001) on the basis of calcareous nannoplankton data, suggests that nearly exclusively syndimentary nannoplankton occurs in the resedimented deposits. This is difficult to comprehend, particularly in typical Magura Beds, which show features indicative of intensive, large-scale resedimentation.

Recycling of dinocysts

Recycling of dinocysts appears to have occurred in the studied sediments on a much larger scale than in coeval sediments in other parts of the flysch Carpathians. Pre-Eocene dinocysts are infrequent, represented mainly by single specimens. The oldest are Middle Jurassic species, represented by *Ctenido-*

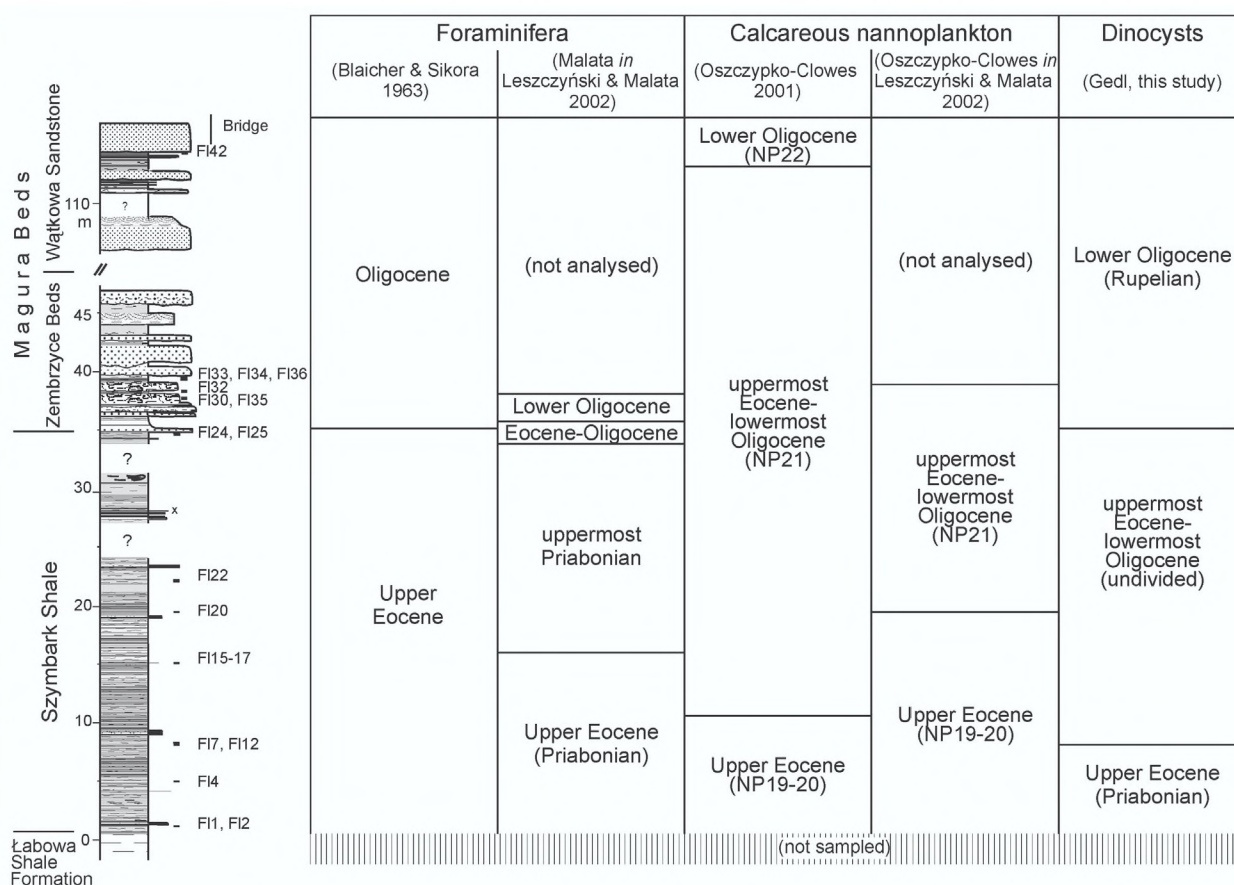


Fig. 7. Bio- and chronostratigraphic interpretation of studied section.

dinium ornatum-*C. combazii*, *Nannoceratopsis gracilis* and *Nannoceratopsis dictyambonis*. They were found in the lowermost part of the Szymbark Shale (samples Fl2 and Fl4) and in the Zembrzyce Beds (samples Fl32 and Fl37). One Early Cretaceous (*Pseudoceratium pelliferum*) and rare Late Cretaceous dinocysts (e.g. *Chatangiella ditissima*, *Trigonopyxidia ginel-la*) were found mainly in the Szymbark Shale. One specimen of Late Paleocene-Early Eocene *Apectodinium quinquelatum* was found in the Zembrzyce Beds (sample Fl37).

The Eocene taxa dominate among the dinocysts believed to be recycled. The most frequent are the Late Eocene species found in the Lower Oligocene part of the section. Mud-flow deposits (samples Fl30 and Fl35) and associated deposits (sample Fl36) exposed in the basal part of the Zembrzyce Beds contain numerous specimens of *Areosphaeridium michoudii*, *Areosphaeridium diktyoplokum* and *Cordosphaeridium funiculatum*. Their presence indicates that these deposits represent Upper or Middle Eocene sediments recycled during the Early Oligocene. Several Middle Eocene species were found in the Upper Eocene part of the section (e.g. *Diphyes pseudoficusoides*), while a single specimen of Early-Middle Eocene *Eatonicysta ursulea* was found in the Oligocene part of the Szymbark Shale.

There is no difference in the state of preservation between the forms treated as recycled or the *in situ* ones. That is why it is difficult to distinguish whether a given species with known early Late Eocene top-range (e.g. *Areosphaeridium michoudii*) is recycled or not in the higher part of the Upper Eocene of the studied section.

Paleoenvironment

The paleoenvironmental conditions in the northern part of the Magura Basin during the Late Eocene and Early Oligocene were reconstructed earlier mainly on the basis of lithofacies distribution and foraminiferal associations (e.g. Książkiewicz Ed. 1962; Sikora 1970; Bromowicz 1992; Leszczyński & Malata 2002). The shape of the basin, its bathymetry and bottom life conditions were the most essential in these interpretations. The palynological, and particularly dinocyst data presented in this paper, supply new information about paleoenvironmental conditions in this part of the Magura Basin.

A characteristic feature of the Eocene-Oligocene transition in the Polish Outer Carpathians is a significant change of the dinocyst assemblage. Diversified Late Eocene dinocyst assemblages disappeared during the earliest Oligocene or they were replaced by Peridinioid-dominated assemblages represented mainly by *Deflandrea* (Fig. 4.19) and *Wetzeliella* (Fig. 4.20). At the same time, a considerable increase of land-derived elements of palynofacies and amorphous organic matter is recorded (Gedl 1999). These changes have been recorded from several sections within the Silesian, Dukla and Skole tectonic units, and from a single section (Leluchów) within the southernmost zone of the Magura Nappe. The changes were interpreted as a result of eutrophication of the surface waters caused by fresh water influx into the Carpathian Flysch Basin (Gedl 1999) and appearance of disoxic/anoxic bottom conditions during sedimentation of the Early Oligocene Menilite Beds (or their equivalents).

The changes that took place during the Eocene-Oligocene transition in the northern part of the Magura Basin seem to differ from those in other parts of the Carpathian Flysch Basin. No such prominent change in the dinocyst assemblages, comparable to that known from coeval sediments of the other parts of the Outer Carpathians, was noted in the studied material. However, this may result from increased reworking, especially in the turbidites. A change in the quality of the dinocyst assemblages is recorded when hemipelagic samples exclusively are taken into account. Among the samples treated as hemipelagic Lower Oligocene sediments (Table 1), two (Fl32 and Fl41) are devoid of dinocysts or contain very infrequent ones. This may result from the appearance of paleoenvironmental conditions within the photic zone, which were inconvenient for dinoflagellates. Noteworthy, palynofacies implies land proximity of the sedimentary area.

Paleogeography: The location of the Siary Zone in marginal part of the Magura Nappe together with lithofacies distribution and paleocurrent directions were long ago interpreted as indicative of location of this zone close to the northern margin of the Magura Basin (e.g. Książkiewicz Ed. 1962). However, the presence of *Impagidinium* spp. in hemipelagites indicates an offshore setting of the sedimentary area of the studied deposits. The lack or infrequent occurrence of this genus in turbidites, where near-shore species prevail, indicates resedimentation of turbidites from inshore areas bordered by a landmass. Presence of a land on this side of the Magura Basin is suggested by the mass occurrence of land plant remains, which dominate the turbidite palynofacies. The domination of terrestrial elements (phytoclasts and sporomorphs) may result from a strong runoff, although scarcity of Peridinales suggests no freshwater influx into this part of the Magura Basin (cf. e.g. Biffi & Grignani 1983). Phytoclasts and sporomorphs could also be recycled from older sediments as it is supported by the presence of recycled dinocysts, obviously derived from older sediments.

A ridge, suggested by Książkiewicz (1956), called the Silesian cordillera (Fig. 8), is generally accepted as the northern border of the Magura Basin. It remains an open question whether it was a land or a submerged ridge. Nemčok et al. (2000) proposed a different paleogeographical setting for the Magura Basin, with the Bohemian Massif as its northern border. The palynological analysis does not solve the question whether the Bohemian Massif or Silesian cordillera was the northern margin of the Magura Basin, but in the latter case, at least some parts of this ridge, at least for some time, must have formed islands covered by vegetation.

A different paleogeographic location is identified for the southernmost part of the Magura Basin during the Late Eocene-Early Oligocene. Late Eocene dinocysts (numerous *Impagidinium* and *Nematosphaeropsis* spp.; see Dale 1996) from coeval deposits at Leluchów (Krynica Zone) imply a depositional setting located far from the seashore (Gedl 2004). The change in the Leluchów dinocyst assemblages and palynofacies close to the Eocene-Oligocene boundary (established on the basis of the highest occurrence of *Areosphaeridium diktyoplokum*) is similar to that known from the outer nappes of the Carpathians. The low-salinity dinocysts (mainly *Deflandrea* spp.) dominate in the lowermost Oligocene of Le-

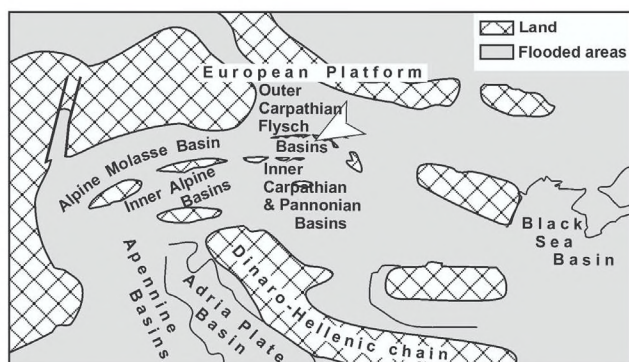


Fig. 8. Schematic paleogeographic map of Alpine and peri-Alpine part of Europe during Eocene-Oligocene transition (based on Rögl 1999, modified by S. Leszczyński). Probable location of investigated section arrowed.

luchów, and disappear higher up the section. These conclusions imply major regional differences between northern and southern margins of the Magura Basin during the Late Eocene and Early Oligocene.

Sea surface temperature: Dinoflagellates, that is inhabitants of the photic zone, as well as their cysts serve as a tenable indicator of the sea surface temperature. The Eocene-Oligocene transition is known as a time of marked global cooling (e.g. Pomeroy & Premoli Silva 1986). Brinkhuis (1994), who traced the records of high latitude dinocyst species within the Eocene-Oligocene boundary interval in Mediterranean, recognized sea surface temperature fluctuations. For this reason he used several, both inshore and offshore species, whereas Zevenboom (1995) argued that only oceanic taxa are suitable for proper estimation of the sea surface temperature. Three species of oceanic dinocysts found in the Folusz section are treated as high latitude ones. These are *Impagidinium velorum* (Fig. 4.6), *Impagidinium pallidum* (Fig. 4.13) and *Gelatia inflata* (Fig. 4.16) described for the first time by Bujak (1984) from the Bering Sea. These species have been found in the lower part of the Szymbark Shale. Their presence may reflect a cooling phase in the Late Eocene, which caused a drop in sea surface temperature during sedimentation of the lower part of the Szymbark Shale. This event seems to correspond to that documented in coeval sediments of the other parts of the Carpathians where high latitude dinocysts *Impagidinium velorum*, *Impagidinium pallidum* and *Gelatia inflata* have been found as well (Gedl unpubl.).

Bottom water oxygenation: Analysis of trace-fossil and benthic foraminiferal assemblages from the Siary Zone indicate poorly oxic to slightly dysoxic conditions prevailing in the bottom waters during the Late Eocene and Early Oligocene (Leszczyński & Malata 2002). This interpretation is supported by the results of our study. The studied sediments contain only occasional traces of amorphous organic matter, which is usually associated with anoxic environments (e.g. Batten 1996). This feature highly contrasts with high content of amorphous organic matter in the Lower Oligocene Menilite Beds and its equivalents in the other parts of the flysch Carpathians (Gedl unpubl.). A better ventilation of the sea floor in the northern part of the Magura Basin during the Early Oli-

gocene might have been caused by intensified water circulation resulting from a high frequency of large-scale resedimentation events (Leszczyński 2001).

Conclusions

1. Dinocysts indicate that the lower part of the Szymbark Shale in the Folusz section (up to the sample Fl7) represents the Upper Eocene (Priabonian). This interpretation is based on continuous presence of *Areosphaeridium michoudii*, *Areosphaeridium diktyoplokum* and *Rhombodinium perforatum*. Several species characteristic for the Middle Eocene (e.g. *Heterelaucacysta porosa*, *Diphyes pseudoficusoides*) that occur in this part of the section are recycled. The uppermost part of the Szymbark Shale, located above the sample Fl24, and the overlying Magura Beds represent the Lower Oligocene. This interpretation is based on the lowest occurrence of *Areoligera? semicirculata* in the sample Fl24. *Wetzeliella gochtii*, a typical Oligocene species in north-western Europe, has the lowest occurrence in the studied section in sample Fl20, dated by foraminifers as Priabonian. Another Oligocene species, *Reticulatosphaera actinocoronata*, has its lowest occurrence in the basal part of the section, dated by other dinocysts, foraminifers and calcareous nannoplankton as Priabonian.

2. Two dinocyst assemblages differing in origin occur in the fine-grained sediments of the entire section. Sediments recognized as redeposited contain frequent near-shore dinocysts. In contrast, the fine-grained sediments interpreted as background deposits always contain specimens of oceanic dinocyst genus *Impagidinium*.

3. No apparent change in quality of dinocyst assemblages is observed in the studied section between the Late Eocene and Early Oligocene assemblages. This contrasts with substantial changes in dinocyst assemblages in coeval sediments of the other parts of the Polish Carpathians. This difference implies various paleoenvironmental conditions, including nutrient availability and salinity level in the photic zone of the northern part of the Magura Basin. However, increased dinocyst recycling observed in the Folusz section may significantly mask the true conditions. Two Lower Oligocene hemipelagic samples containing poor dinocyst assemblages appear to reflect inconvenient paleoenvironmental conditions in the photic zone.

4. The appearance of high-latitude *Impagidinium velorum*, *Impagidinium pallidum* and *Gelatia inflata* in the Upper Eocene of the investigated deposits is interpreted as resulting from a drop of sea surface temperature in the Magura Basin during the latest Eocene. This event, which is also recorded in coeval sediments of the other parts of the Polish Carpathians, seems to be of regional significance and appear to correspond to the global cooling recorded across the Eocene-Oligocene boundary.

Acknowledgments: We thank H. Brinkhuis, J. P. Bujak and J. Soták for their critical remarks that improved the manuscript greatly. The Folusz section was described and sampled by Stanisław Leszczyński in the frame of the project financed by the State Committee for Scientific Research (Komitet Badań

Naukowych) research Grant No. 6 P04D 021 16. Palynological study was done by Przemysław Gedl as a part of the State Committee for Scientific Research (Komitet Badań Naukowych) research Grant No. 6P04D 042 15.

References

- Batten D.J. 1996: Palynofacies and palaeoenvironmental interpretation. In: Jansonius J. & McGregor D.C. (Eds.): *Palynology: principles and applications. Amer. Assoc. Stratigraphic Palynologist Foundation*, Dallas, Texas, 3, 1011–1064.
- Benedek P.N. 1986: Ergebnisse der Phytoplankton-Untersuchungen aus dem Nordwestdeutschen Tertiär. In: Tobien H. (Ed.): *Nordwestdeutschland im Tertiär*. Berlin, 157–185.
- Berggren W.A., Kent D.V., Swisher C.C. & Aubry M.-P. 1995: A revised Cenozoic geochronology and chronostratigraphy. In: Berggren W.A., Kent D.V. & Hardenbol J. (Eds.): *Geochronology, Time Scales and Global Stratigraphic Correlation. Society of Economic Paleontologists and Mineralogists, Spec. Publ.* 54, 129–212.
- Biffi U. & Grignani D. 1983: Peridinioid dinoflagellate cysts from the Oligocene of the Niger Delta, Nigeria. *Micropaleontology* 29, 126–145.
- Biffi U. & Manum S.B. 1988: Late Eocene-Early Miocene dinoflagellate cyst stratigraphy from the Marche Region (central Italy). *Boll. Soc. Paleont. Ital.* 27, 163–212.
- Birkenmajer K. & Oszczyk N. 1989: Cretaceous and Paleogene lithostratigraphic units of the Magura Nappe, Krynica Subunit, Carpathians. *Ann. Soc. Geol. Pol.* 59, 145–181.
- Blaicher J. & Sikora W. 1963: Attempt to correlate the Magura beds in eastern part of Magura nappe with sediments of external group. *Kwart. Geol.* 7, 620–626 (in Polish with English summary).
- Brinkhuis H. 1994: Late Eocene to Early Oligocene dinoflagellate cysts from the Priabonian type-area (northeast Italy); biostratigraphy and paleoenvironmental interpretation. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 107, 121–163.
- Brinkhuis H. & Biffi U. 1993: Dinoflagellate cyst stratigraphy of the Eocene/Oligocene transition in central Italy. *Mar. Micropaleontology* 22, 131–183.
- Brinkhuis H. & Visscher H. 1995: The upper boundary of the Priabonian Stage; a reappraisal based on dinoflagellate cyst biostratigraphy. In: Berggren W.A., Kent D.V. & Hardenbol J. (Eds.): *Geochronology, Time Scales and Global Stratigraphic Correlation. Society of Economic Paleontologists and Mineralogists, Spec. Publ.* 54, 295–304.
- Bromowicz J. 1992: The sedimentary basin and source areas of the Magura Sandstones. *Zesz. Nauk. Akad. Gór.-Hutn., Geol.* 54, 1–116 (in Polish with English summary).
- Bujak J.P. 1984: Cenozoic dinoflagellate cysts and acritarchs from the Bering Sea and northern North Pacific, D.S.D.P. Leg 19. *Micropaleontology* 30, 180–212.
- Bujak J.P. & Mudge D. 1994: A high-resolution North Sea Eocene dinocyst zonation. *J. Geol. Soc., London* 151, 449–462.
- Châteauneuf J.-J. 1980: Palynostratigraphie et paléoclimatologie de l'Eocène supérieur et de l'Oligocène du Bassin de Paris (France). *Mém. Bur. Rech. Géol. Min.* 116, 1–360.
- Coccioni R., Basso D., Brinkhuis H., Galeotti S., Gardin S., Monechi S. & Spezzaferri S. 2000: Marine biotic signals across a late Eocene impact layer at Massignano, Italy: evidence for long-term environmental perturbations? *Terra Nova* 12, 258–263.
- Costa L.I. & Downie C. 1976: The distribution of the dinoflagellate *Wetzeliiella* in the Palaeogene of north-western Europe. *Palaeontology* 19, 591–614.
- Dale B. 1996: Dinoflagellate cyst ecology: modelling and geological applications. In: Jansonius J. & McGregor D.C. (Eds.): *Palynology: principles and applications. Amer. Assoc. Stratigraphic Palynologist Foundation*, Dallas, Texas, 3, 1249–1275.
- El-Beialy S.Y. 1988: The dinocyst biostratigraphy of the Upper Eocene subsurface sediments, west Nile Delta, Egypt. *Newslett. Stratigr.* 19, 131–141.
- Gedl P. 1995: Middle Eocene dinoflagellate cysts from the Rogoźnik section, Flysch Carpathians, Poland. *Acta Palaeobot.* 35, 195–231.
- Gedl P. 1999: Palynology of the Eocene-Oligocene boundary in the Polish Flysch Carpathians. *Przegl. Geol.* 47, 394–399 (in Polish with English summary).
- Gedl P. 2000: Biostratigraphy and palaeoenvironment of the Podhale Paleogene (Inner Carpathians, Poland) in the light of palynological studies. Part II. Summary and systematic descriptions. *Stud. Geol. Pol.* 117, 155–303.
- Gedl P. 2004: Dinoflagellate cyst record of the Eocene-Oligocene boundary succession in flysch deposits at Leluchów, Carpathian Mountains, Poland. *Geol. Soc. Spec. Publ., London* 230, 309–324.
- Gruas-Cavagnetto C. & Barbin V. 1988: Dinoflagellates from the Priabonian stage stratotype area (Vicentin, Italy), evidence for the Eocene/Oligocene boundary. *Rev. Paléobiologie* 7, 163–198 (in French with English summary).
- Kopciowski R. 1996: Geological structure of the Siary zone between Ropa and Banica (the Magura Nappe). *Biul. Państw. Inst. Geol.* 374, 21–40 (in Polish with English summary).
- Koszarski L. 1976: Development of the youngest sediments in the eastern part of the Magura Nappe. *Spraw. Posiedz. Kom. Nauk. PAN Krakow* 20, 174–175 (in Polish).
- Koszarski L. & Tokarski A. 1968: Detailed Geological Map of Poland (without Quaternary deposits). Sheet M34-91B Osiek — provisional edition. *Inst. Geol., Warszawa* (in Polish).
- Koszarski L. & Wieser T. 1960: New tuff horizons in older Palaeogene of Flysch Carpathians. *Kwart. Geol.* 4, 749–771 (in Polish with English summary).
- Koszarski L., Sikora W. & Wdowiarz S. 1974: The Flysch Carpathians. Polish Carpathians. In: Mahel' M. (Ed.): *Tectonics of the Carpathian-Balkan Regions. Geologický Ústav Dionýza Štúra*, Bratislava, 180–197.
- Książkiewicz M. 1956: Geology of the northern Carpathians. *Geol. Rdsch.* 45, 369–411.
- Książkiewicz M. (Ed.) 1962: Geological Atlas of Poland. Stratigraphic and facial problems. Fascicle 13, Cretaceous and Early Tertiary in the Polish External Carpathians. *Inst. Geol., Warszawa*.
- Książkiewicz M. 1974: Explanations to the Detailed Geological Map of Poland. Sheet Sucha Beskidzka (1014), 1:50,000. *Wydawnictwa Geologiczne* 1–83 (in Polish).
- Książkiewicz M. 1977: The tectonics of the Carpathians. In: Książkiewicz M. (Ed.): *Geology of Poland, IV, Tectonics. Wydawnictwa Geologiczne*, Warszawa, 476–618.
- Leszczyński S. 2001: Carpathian flysch sea in Late Eocene-Early Oligocene: example of oxygenation decline. *IAS 2001 21st Meeting, Davos, Switzerland, 3–5. September 2001, Abstracts & Programme*, 99–100.
- Leszczyński S. & Malata E. 2002: Sedimentary conditions in the Siary zone of the Magura basin (Carpathians) in the Late Eocene-Early Oligocene. *Ann. Soc. Geol. Pol.* 72, 201–239.
- Maier D. 1959: Systematik, Stratigraphie und Ökologie der Coccolithophorideen, Dinoflagellaten und Hystrichosphaerideen vom Oligozän bis Pleistozän. *Neu. Jb. Geol. Paläont., Abh.* 107, 278–340.
- Manum S.B., Boulter M.C., Gunnarsdottir H., Rangnes K. & Scholze A. 1989: Eocene to Miocene palynology of the Norwe-

- gian Sea (ODP Leg 104). In: Eldholm O., Thiede J., Taylor E. et al. *Proceedings of the Ocean Drilling Program, Scientific Results*, 104, 611–662.
- Martini E. 1971: Standard Tertiary and Quaternary calcareous nannoplankton zonation. In: Farinacci A. (Ed.): *Proceedings of the 2nd Planktonic Conference, Roma, 1970. Edizioni Technoscienza, Roma*, 739–785.
- Morgenroth P. 1966: Neue in organischer Substanz erhaltene Mikrofossilien des Oligozäns. *Neu. Jb. Geol. Paläont., Abh.* 127, 1–12.
- Nemčok M., Nemčok J., Wojtaszek M., Ludhová L., Klecker R.A., Sercombe W.J., Coward M.P. & Keith J.F. Jr. 2000: Results of 2D balancing along 2030 longitude and pseudo-3D in the Smilno tectonic window: implications for shortening mechanisms of the West Carpathian accretionary wedge. *Geol. Carpathica* 51, 281–300.
- Olszewska B. 1984: Benthonic foraminifera of the Sub-Menilite Globigerina Marls of Polish Outer Carpathians. *Prace Inst. Geol.* 110, 1–37 (in Polish with English summary).
- Olszewska B. 1985: Remarks concerning the Eocene-Oligocene boundary in the Polish External Carpathians: results of foraminiferal investigations. *Proceedings reports of the XIII-th Congress of KBGA, Poland – Cracow, 5.–10. September 1985*. 1, 57–59.
- Oszczypko N. 1991: Stratigraphy of the Paleogene Deposits of the Bystrica Subunit (Magura Nappe, Polish Outer Carpathians). *Bull. Pol. Acad. Sci., Earth Sci.* 39, 415–431.
- Oszczypko-Clowes M. 2001: The nannofossil biostratigraphy of the youngest deposits of the Magura Nappe (east of the Skawa river, Polish Flysch Carpathians) and their paleoenvironmental conditions. *Ann. Soc. Geol. Pol.* 71, 139–188.
- Pomerol C. & Premoli Silva I. 1986: The Eocene-Oligocene transition: events and boundary. In: Pomerol C. & Premoli Silva I. (Eds.): *Terminal Eocene Events. Developments in Palaeontology and Stratigraphy. Elsevier, Amsterdam*, 9, 1–24.
- Powell A.J. 1992: Dinoflagellate cysts of the Tertiary System. In: Powell A.J. (Ed.): *A stratigraphic index of dinoflagellate cysts. Brit. Micropalaeontological Soc. Publ. Ser., Chapman & Hall, London*, 155–249.
- Rögl F. 1999: Mediterranean and Paratethys. Facts and hypotheses of an Oligocene to Miocene paleogeography (short overview). *Geol. Carpathica*, 50, 339–349.
- Sikora W. 1970: Geology of the Magura nappe between Szymbark Ruski and Nawojowa. *Inst. Geol., Biul.* 235, 1–97 (in Polish with English summary).
- Stover L.E. & Hardenbol J. 1993: Dinoflagellates and depositional sequences in the Lower Oligocene (Rupelian) Boom Clay Formation, Belgium. *Bull. Soc. Belge Géol., Bulletin van de Belgische Vereniging voor Geologie* 102, 5–77.
- Stover L.E., Brinkhuis H., Damassa S.P., de Verteuil L., Helby R.J., Monteil E., Partridge A.D., Powell A.J., Riding J.B., Smelror M. & Williams G.L. 1996: Mesozoic-Tertiary dinoflagellates, acritarchs and prasinophytes. In: Jansonius J. & McGregor D.C. (Eds.): *Palynology: principles and applications. Amer. Assoc. Stratigraphic Palynologist Foundation, Dallas, Texas*, 2, 641–750.
- Szajnocha W. 1895: Geological Atlas of Galicyja. Text to the sixth volume. *Wydział Krajowy Królestwa Galicyji i Lodomeryji*, 1–149 (in Polish).
- Świdziński H. 1934: Recherches géologiques dans les environs de Wysowa-les-Bains, district Gorlice, Carpathes. *Posiedz. Nauk. Państw. Inst. Geol.* 38, 29–30.
- Van Couvering J.A., Aubry M.-P., Berggren W.A., Bujak J.P., Naeser C.W. & Wieser T. 1981: The terminal Eocene event and the Polish connections. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 36, 321–362.
- Van Mourik C.A. & Brinkhuis H. 2000: Data report: Organic walled dinoflagellate cyst biostratigraphy of the latest middle to late Eocene at Hole 1053 (subtropical Atlantic Ocean). In: Kroon D., Norris R.D. & Klaus A. (Eds.): *Proceedings of the Ocean Drilling Project, Scientific Results*. 171B, 1–25.
- Wilpshaar M., Santarelli A., Brinkhuis H. & Visscher H. 1996: Dinoflagellate cysts and mid-Oligocene chronostratigraphy in the central Mediterranean region. *J. Geol. Soc., London* 153, 553–561.
- Zevenboom D. 1995: Dinoflagellate cysts from the Mediterranean Late Oligocene and Miocene. *Ph. D. Thesis, University of Utrecht*, 1–221.